AMENDMENTS TO THE CLAIMS

This listing of claims replaces all prior versions, and listings, of claims in the application.

1. (Currently Amended) A pharmaceutical composition, ~~comprised of~~ comprising a non-steroidal anti-inflammatory and an opiate analgesic in combination with colloidal silicate dioxide, sodium glycolate ~~glicolate~~, lactose, microcrystalline cellulose, and magnesium stearate ~~estearate and other recipients if necessary~~.

2. (Original) A pharmaceutical composition according to claim 1, wherein the non-steroidal anti-inflammatory is ketorolac tromethamine and the opiate analgesic is tramadol hydrochloride.

3. (Currently Amended) A pharmaceutical composition according to claim ~~1~~ 2, wherein the ketorolac tromethamine is present in the composition in a range of 0.0010 g to 0.1000 g and the tramadol hydrochloride is present in a range of 0.0010 g to 0.2000 g.

4. (Currently Amended) A pharmaceutical composition according to claim ~~1~~ 2, wherein the ketorolac ~~ketorolace~~ tromethamine is present in the composition in a range of 0.0010 g to 0.10000 g and the tramadol hydrochloride is present at a range of 0.0010 g to 0.2000 g, the colloidal silicate dioxide is present in a range of 0.00010g to 0.02000g, the sodium glycolate ~~glicolate~~ starch is present in a range of 0.0010g to 0.20000g, the lactose is present in the range of 0.0100g to 0.50000g, the microcrystalline cellulose is present in a range of 0.0100g to 0.50000g, the magnesium stearate is present at a proportion of 0.0001g to 0.02000g, ~~other recipients~~ and the excipient is ~~can be~~ present at a proportion of 0.0001g to 1.000g.

5. (Original) A pharmaceutical composition according to claim 1, wherein the composition ~~it~~ is in a capsule form.

6. (Currently Amended) A method for preparing ~~procedure for the elaboration of~~ a composition according to claim ~~1~~ 2, ~~wherein it is comprised of~~ comprising the following steps:

a) ~~Mix~~ mixing the ketorolac tromethanine, ~~the~~ colloidal silicate dioxide, ~~the~~ tramadol hydrochloride, sodium glycolate ~~glicolate starch~~, ~~the~~ lactose, ~~the~~ microcrystalline cellulose, ~~the~~ and magnesium stearate to produce a powdered mixture ~~and other recipients if necessary~~;

b) ~~analyze~~ analyzing the powdered ~~mix~~ mixture; and

c) ~~proceed to~~ encapsuling and conditioning the ~~mix~~ mixture.

7. (Currently Amended) ~~The use of~~ A method of treating pain, the method comprising administering an effective amount of the composition according to any one of claims 1 to 6~~, for the pain treatment~~.